K003683

JAN 11 2001

21.0 510(K) SUMMARY

Submitter: **Jeneric/Pentron, Inc.**
Address: **53 North Plains Industrial Road**
Wallingford, Connecticut 06492
Contact Tel: 203-265-7397 X619
Contact Fax: 203-265-7662
Contact Person: Annmarie Tenero

Date Summary Prepared: November 28, 2000

The Avanté Cure Light is a gun-handle shaped, dental curing light that is intended to polymerize dental materials, such as resins and sealants by transmitting light through a light guiding tip. The Avanté Cure Light has two optional curing modes.

The Avanté Cure Light is substantially equivalent to Coltolux 3, K945698, Coltene/Whaledent. The Avanté Cure Light has three selectable time periods and a single or step power cycle. The Coltolux 3, has six selectable time periods and a single output power on all cycles. Safety and effectiveness are not affected because the intended use of the Avanté Cure light is virtually identical to the predicate device. There are only minor differences in devices are setting time periods and power cycles.

Jeneric/Pentron, by its request that FDA find this product to be substantially equivalent, does not waive any rights that the Avanté Cure Light™ product is patentable and further makes no admission that the product infringes any patent covering any of the competitive products mentioned herein.

DEPARTMENT OF HEALTH & HUMAN SERVICES

Public Health Service

Food and Drug Administration
9200 Corporate Boulevard
Rockville MD 20850

JAN 11 2001

Ms. Annmarie Tenero
Jeneric/Pentron, Incorporated
53 North Plains Industrial Road
P.O. Box 724
Wallingford, Connecticut 06492-0724

Re: K003683
Trade Name: Avante Cure Light
Regulatory Class: II
Product Code: EBZ
Dated: November 28, 2000
Received: November 29, 2000

Dear Ms. Tenero:

We have reviewed your Section 510(k) notification of intent to market the device referenced above and we have determined the device is substantially equivalent (for the indications for use stated in the enclosure) to legally marketed predicate devices marketed in interstate commerce prior to May 28, 1976, the enactment date of the Medical Device Amendments, or to devices that have been reclassified in accordance with the provisions of the Federal Food, Drug, and Cosmetic Act (Act). You may, therefore, market the device, subject to the general controls provisions of the Act. The general controls provisions of the Act include requirements for annual registration, listing of devices, good manufacturing practice, labeling, and prohibitions against misbranding and adulteration.

If your device is classified (see above) into either class II (Special Controls) or class III (Premarket Approval), it may be subject to such additional controls. Existing major regulations affecting your device can be found in the Code of Federal Regulations, Title 21, Parts 800 to 895. A substantially equivalent determination assumes compliance with the Current Good Manufacturing Practice requirements, as set forth in the Quality System Regulation (QS) for Medical Devices: General regulation (21 CFR Part 820) and that, through periodic QS inspections, the Food and Drug Administration (FDA) will verify such assumptions. Failure to comply with the GMP regulation may result in regulatory action. In addition, FDA may publish further announcements concerning your device in the Federal Register. Please note: this response to your premarket notification submission does not affect any obligation you might have under sections 531

through 542 of the Act for devices under the Electronic Product Radiation Control provisions, or other Federal laws or regulations.

This letter will allow you to begin marketing your device as described in your 510(k) premarket notification. The FDA finding of substantial equivalence of your device to a legally marketed predicate device results in a classification for your device and thus, permits your device to proceed to the market.

If you desire specific advice for your device on our labeling regulation (21 CFR Part 801 and additionally 809.10 for in vitro diagnostic devices), please contact the Office of Compliance at (301) 594-4692. Additionally, for questions on the promotion and advertising of your device, please contact the Office of Compliance at (301) 594-4639. Also, please note the regulation entitled, "Misbranding by reference to premarket notification" (21CFR 807.97). Other general information on your responsibilities under the Act may be obtained from the Division of Small Manufacturers Assistance at its toll-free number (800) 638-2041 or (301) 443-6597 or at its internet address "http://www.fda.gov/cdrh/dsma/dsmamain.html".

Sincerely yours,

Timothy A. Ulatowski
Director
Division of Dental, Infection Control
and General Hospital Devices
Office of Device Evaluation
Center for Devices and
Radiological Health

Enclosure

5.0 INDICATION FOR USE STATEMENT

510(k) NUMBER (IF KNOWN): K003683

DEVICE NAME: Avanté Cure Light

INDICATION FOR USE: The Avanté Cure Light is a gun-handle shaped, dental curing light that is intended to polymerize dental materials, such as resins and sealants by transmitting light through a light guiding tip.

(PLEASE DO NOT WRITE BELOW THIS LINE – CONTINUE ON ANOTHER PAGE IF NEEDED.)

Susan Perrer

(Division Sign-Off)
Division of Dental, Infection Control, and General Hospital Devices
510(k) Number K003683

Concurrence of CDRH, Office of Device Evaluation (ODE)

Prescription Use ✓ **OR** **Over –The-Counter-Use** _____
(Per 21 CFR 801.109) **(Optional Format 1-2-96)**